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54 Sustained release ibuprofen formulation.

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Description

The present invention relates to a sustained release ibuprofen formulation, and in particular to such a formulation which will provide sustained release of the ibuprofen over a period of about twelve hours.

It is well known that some medical conditions are best treated by administration of a pharmaceutical which is formulated to allow the pharmaceutical to act as quickly as possible. Such a formulation may comprise an injectable solution or a readily dissolvable tablet or capsule. Such formulations are useful, for instance, for treating acute pain, such as headaches or pain associated with trauma, such as an accident.

On the other hand some medical conditions are best treated by administration of a pharmaceutical in such a way as to sustain its action over an extended period of time. This type of administration is useful, for instance, for treating chronic pain, such as that associated with rheumatic or arthritic conditions.

This type of administration can be achieved by repeated administration of an immediate-release tablet or capsule at frequent intervals, for instance every four hours. However, this is generally inconvenient, especially during the night, when it is often necessary to wake a patient up to administer the tablet or capsule.

It has therefore been proposed to produce a formulation which will release the pharmaceutical therein at a controlled rate such that the amount of the pharmaceutical available in the body to treat the condition is maintained at a relatively constant level over an extended period. A particularly suitable period is twelve hours, since such a formulation need only be taken twice a day to maintain control of the condition. Such formulations are generally known as sustained release formulations.

Although many sustained release formulations are known, there is no generally applicable method by which such formulations can be designed. Each formulation is dependent on the particular pharmaceutical incorporated therein. In designing a formulation, it is generally necessary to take into account many factors, including the rates of absorption and clearance of the pharmaceutical, the interaction of the pharmaceutical with the excipients and/or coatings to be used in the formulation, the solubility of the excipients and/or coatings, and the effects on the bioavailability of the pharmaceutical which may be caused by the excipients and/or coatings. It is, however, not possible readily to predict whether any particular formulation will provide the desired sustained release, and it is generally found necessary to carry out considerable experimentation to produce a desired sustained release formulation.

It is known that ibuprofen (which is the generic name for α -methyl-4-(2-methylpropyl)benzene acetic acid; see GB-A-0 971 700) is a pharmaceutical useful for the treatment of both acute and chronic pain. For instance, in the United

Kingdom it is sold in an immediate release formulation as Brufen, Apsifen, Proflex or Nurofen and as a sustained release formulation as Fenbid. (Brufen, Apsifen, Nurofen and Fenbid are all registered trade marks).

It has been found that the sustained release formulation which is on the market does not provide as good a sustained release over a twelve hour period as is desirable. There is therefore a need to provide an improved sustained release ibuprofen formulation having good sustained release over a period of twelve hours. The present invention is based on the discovery of such an improved sustained release ibuprofen formulation.

In the following description, all parts and percentages are by weight unless otherwise indicated.

According to the present invention, there is provided a sustained release ibuprofen formulation comprising sufficient granules to provide a predetermined dose or number of doses of ibuprofen, each granule having a diameter between 0.5 and 2.0 mm and comprising:

a) a core containing 100 parts by wt. of ibuprofen and from 10 to 70 parts by wt. of a microcrystalline cellulose; and

b) a coating covering substantially the whole surface of the core and comprising 100 parts by wt. of a water insoluble but water swellable acrylic polymer and 20 to 70 parts by wt. of a water-soluble hydroxylated cellulose derivative, the weight of the coating being from 5 to 20% of the weight of the core.

It will be appreciated that the exact diameter of the granules and the composition and amount of coating will depend on the time over which the formulation is designed to work.

For instance, for a 24 hour formulation, the granule diameter is preferably between 1.5 and 2.0 mm, the coating preferably contains 25 to 35 parts by wt. of the water soluble hydroxylated cellulose derivative, and the weight of the coating is approximately 20% of the weight of the core.

In another instance, for a 12 hour formulation, the granule diameter is preferably between 0.7 and 1.2 mm, the coating preferably contains 25 to 35 parts by wt. of the water soluble hydroxylated cellulose derivative, and the weight of the coating is between 8 and 12% of the weight of the core.

The core preferably contains between 15 and 25 parts by wt., advantageously 20 parts by wt. of the microcrystalline cellulose, and may conveniently be prepared by mixing the components together with some water to produce a slightly cohesive product. The cohesive product may then be extruded, chopped into suitable lengths, spheronised and dried.

Microcrystalline cellulose is a well known form of cellulose which is partially depolymerised. A particularly suitable microcrystalline cellulose is sold under the name of Avicel (which is a registered trade mark). However, other equivalent materials may be used, as will be readily apparent to those skilled in the art.

Preferably, the coating comprises about 30 parts by wt. of the hydroxylated cellulose derivative. If too much of this derivative is present, the coating may become too sticky and the rate of release may become too high. If too little is present, the rate of release may be too low.

A particularly suitable hydroxylated cellulose derivative is hydroxypropyl methylcellulose having a degree of substitution of 28 to 30% of methoxy groups and 7 to 12% of hydroxy groups. However, other equivalent materials such as hydroxyethyl or hydroxymethyl celluloses can be used.

The acrylic polymer is preferably neutral and may comprise a homopolymer or a copolymer, for instance of acrylic acid esters or methacrylic acid esters. Preferably, the acrylic polymer is provided as an aqueous dispersion.

A particularly suitable acrylic polymer is sold under the name of Eudragit (which is a registered trade mark), which comprises a copolymer of acrylic and methacrylic acid esters and which is usually supplied as an aqueous dispersion containing approximately 30% solids.

The coating may conveniently be prepared by forming a solution of the hydroxylated cellulose derivative and mixing it with a dispersion of the acrylic polymer. The aqueous mixture is then used to coat the dried core particles, and the coated particles are subsequently dried to produce the granules.

Preferably, the coated granules are sieved to ensure that they are in the correct size range.

The granules may be supplied loose with a means for dispensing a measured amount of granules, for instance to be sprinkled on food. Alternatively, the granules may be provided in sachets containing measured amounts.

The granules may, if desired, be formed into tablets using conventional tableting machinery.

However, preferably the granules are placed in measured amounts in readily soluble capsules. The capsule may be any of those already known in the art, and may, for instance, comprise a thin gelatin skin. Preferably, the capsule contains either 200 or 300 mg of Ibuprofen.

It has surprisingly been found that by the above set out selection of the materials for the core and the coating, the relative amounts of the components, and the size of the granules, it has proved possible to produce a sustained release Ibuprofen formulation which acts effectively over any desired period, in particular a twelve hour period.

One embodiment of the present invention is now described, by way of example only, with reference to the accompanying drawings, in which:

Figure 1 shows the mean plasma concentration of Ibuprofen following administration of Brufen;

Figure 2 is similar to Figure 1, but relates to Fenbid;

Figure 3 is similar to Figure 1, but relates to a formulation according to the present invention; and

Figure 4 shows blood level time profiles after a fourth day morning dose following three days of administration of a formulation according to the invention (—O—) and Brufen (—Δ—).

Example

The formulation described below was developed using the OSAT system developed by the Inventors at the University of Bradford.

1000 g of Ibuprofen and 200 g of Avicel were mixed together by doubling up in a dry blender. 615 ml of water was added in portions until a slightly cohesive product was formed. The cohesive product was passed through an extruder and the extruded material was chopped to produce slugs having a diameter of about 1 mm and a length of 2 to 3 mm. The slugs were spheronised by passage through a spheroniser, and the particles thus formed were dried to constant weight at 45°C. The dried particles were sieved to separate those having diameters between 0.7 and 1.5 mm.

15 g of hydroxypropyl methylcellulose was dissolved in 135 ml of hot water and cooled. The cooled solution was mixed with 167 g of Eudragit E30D (containing 50 g solids) and the mixture was diluted with a further 190 ml of water to produce a coating mixture containing 0.128 g solids per gram of mixture.

The sieved core particles were rotated in a small coating pan and the coating mixture was added in portions to the pan until the weight of solids in the added coating mixture was 10% of the weight of the core particles. After each portionwise addition of coating mixture, air was blown into the pan to assist in solvent removal. At the end of the addition of the coating mixture, the coated core particles were dried to constant weight at ambient temperature and sieved to produce granules having a size between 0.8 and 1.2 mm.

Hard gelatin capsules were each filled with 400 mg of the granules, to produce a total dose of 300 mg of Ibuprofen per capsule.

The following experiments were carried out on healthy human volunteers to determine the release properties of the formulation set out above in comparison to Brufen and Fenbid.

Each volunteer had a light evening meal and drank 1.5 l of tap water during the course of the evening prior to each trial.

For the trials with Brufen, which is an immediate release formulation, each volunteer had a light breakfast at 08.00 hrs and took an oral dose of 200 mg Brufen at 10.30 hrs. Blood samples were collected at 0.5, 1, 1.5, 2, 4, 6, 8 and 12 hours after administration of the dose.

For the trials with Fenbid and the formulation of the invention, each volunteer took an oral dose of 300 mg of Ibuprofen, either as Fenbid or as formulated above, at 07.30 hrs and then had a light breakfast at 09.00 hrs. Blood samples were collected at 0, 2, 4, 6, 8, 10, 12 and 15 hours after administration of the dose.

During all the trials, only light meals were taken

and each volunteer was limited to a daily fluid intake of approximately 1.5 l of water.

A washout period of one week was allowed between each of the three series of trials.

Blood samples were collected, by finger tip puncture with a sterile blood lancet, into heparinized capillary collection tubes. Plasma was harvested by centrifugation and deep-frozen until assayed. Assays were carried out by high pressure liquid chromatography on a 250x4.6 mm reversed phase column packed with Partisil 10 ODS at ambient temperature, using a mobile phase of 50% acetonitrile/50% water at pH 3, using a solution of butyl-p-hydroxybenzoate as an internal standard.

Referring now to the Figures, it can be seen from Figure 1 that the plasma level of Ibuprofen following administration of Brufen rises rapidly to a peak at about 2 hours and rapidly falls to a low level.

From Figure 2 it can be seen that for Fenbid the peak value occurs at approximately 4 hours with a less rapid rise and fall than for Brufen. However, the plasma level is by no means as constant as desirable.

From Figure 3 it can be seen that the plasma level obtained using the present formulation is much more constant than that obtained even with Fenbid, the level being relatively constant over a period of at least 2 to 12 hours following administration.

A measure of the sustained release properties of a formulation is the ratio between the plasma level at its peak and the plasma level at the end of the desired sustained release period, in this case 12 hours. This ratio preferably is as close to one as possible. For Brufen this ratio is 18.4, for Fenbid it is 5.3 and for the formulation of the present invention it is 2.1.

It can readily be seen from the data presented above that the present formulation provides a much improved twelve hour sustained release formulation than was previously available.

A further advantage of the present formulation is that it can be taken with or without food. If the formulation is taken with food, the time taken to reach its peak concentration in plasma is about the same as if it is taken without food, whereas the peak plasma concentration is slightly reduced and the absorption is spread over a slightly longer period. However, these differences are only minor and can for practical purposes be ignored, as the usual statistical tests did not show them to be significant.

The present formulation has also been compared with the immediate-release formulation Brufen in a multi-dose, double-blind crossover study using 12 healthy human volunteers. Figure 4 summarises the results obtained. The study confirmed the pronounced sustained release characteristics of the present formulation and provided clear evidence of persisting drug levels in plasma in the morning following an evening dose, a phenomenon which is much less pronounced (and totally absent in some subjects)

when Brufen is used. Therefore, the present formulation has the potential to relieve "morning stiffness", a common problem in rheumatic and arthritic patients. Also, the formulation has substantially no effect on the bioavailability of the Ibuprofen compared with the known immediate-release formulations such as Brufen and Apsifen, although the apparent bioavailability of both was somewhat lower in the multi-dose study than in the single-dose studies.

Moreover, it has been shown that the formulation is storage stable over at least 12 months at high temperatures (up to 50°C) and high relative humidity. There is little effect on the Ibuprofen or the sustained release properties under these stressful storage conditions and no effect under normal storage conditions.

Claims

1. A sustained release α -methyl-4-(2-methylpropyl)benzene acetic acid (Ibuprofen) formulation comprising sufficient granules to provide a predetermined dose or number of doses of Ibuprofen, each granule having a diameter between 0.5 and 2.0 mm and comprising:

a) a core containing 100 parts by wt. of Ibuprofen and from 10 to 70 parts by wt. of a microcrystalline cellulose; and

b) a coating covering substantially the whole surface of the core and comprising 100 parts by wt. of a water insoluble but water swellable acrylic polymer and 20 to 70 parts by wt. of a water-soluble hydroxylated cellulose derivative, the weight of the coating being from 5 to 20% of the weight of the core.

2. The formulation of claim 1, which is a 24 hour formulation, wherein the granule diameter is between 1.5 and 2.0 mm, the coating contains 25 to 35 parts by wt. of the water soluble hydroxylated cellulose derivative, and the weight of the coating is approximately 20% of the weight of the core.

3. The formulation of claim 1, which is a 12 hour formulation, wherein the granule diameter is between 0.7 and 1.2 mm, the coating contains 25 to 35 parts by wt. of the water soluble hydroxylated cellulose derivative, and the weight of the coating is between 8 and 12% of the weight of the core.

4. The formulation of any one of claims 1 to 3, wherein the core contains between 15 and 25 parts by wt. of the microcrystalline cellulose.

5. The formulation of any one of claims 1 to 4, wherein the coating comprises about 30 parts by wt. of the hydroxylated cellulose derivative.

6. The formulation of any one of claims 1 to 5, wherein the hydroxylated cellulose derivative is hydroxypropyl methylcellulose having a degree of substitution of 28 to 30% of methoxy groups and 7 to 12% of hydroxy groups.

7. The formulation of any one of claims 1 to 6, wherein the acrylic polymer is neutral.

8. The formulation of any one of claims 1 to 7, wherein the granules are contained in a capsule.

9. A process for producing a formulation according to any one of claims 1 to 8 comprising:

1. mixing the ibuprofen with the microcrystalline cellulose;
2. forming the mixture into particles;
3. forming a solution of the hydroxylated cellulose derivative and the acrylic polymer; and
4. coating the particles using the solution to form the granules.

10. The process of claim 9, wherein the granules are sieved after formation.

Patentansprüche

1. α -Methyl-4-(2-methylpropyl)benzol-Essigsäure-(Ibu-profen-) Formulierung mit verzögerter Wirkstoffabgabe, die genug Körnchen enthält, um eine vorherbestimmte Dosis oder Zahl von Dosen Ibuprofen zu bieten, wobei jedes Körnchen einen Durchmesser zwischen 0,5 und 2,0 mm aufweist und

a) einen Kern, der 100 Gewichtsteile Ibuprofen und 10 bis 70 Gewichtsteile einer mikrokristallinen Zellulose enthält; und

b) eine Umhüllung umfaßt, die im wesentlichen die gesamte Oberfläche des Kerns bedeckt und 100 Gewichtsteile eines wasserunlöslichen aber in Wasser schwellenden Akrylpolymeren und 20 bis 70 Gewichtsteile eines wasserlöslichen, hydroxylierten Zellulosederivates enthält, wobei das Gewicht der Umhüllung 5 bis 20% des Kerngewichtes beträgt.

2. Formulierung nach Anspruch 1, die eine 24-Stunden-Formulierung ist, wobei der Körnchendurchmesser 1,5 bis 2,0 mm beträgt, die Umhüllung 25 bis 35 Gewichtsteile des wasserlöslichen, hydroxylierten Zellulosederivates enthält und das Gewicht der Umhüllung etwa 20% des Kerngewichtes beträgt.

3. Formulierung nach Anspruch 1, die eine 12-Stunden-Formulierung ist, wobei der Körnchendurchmesser 0,7 bis 1,2 mm beträgt, die Umhüllung 25 bis 35 Gewichtsteile des wasserlöslichen, hydroxylierten Zellulosederivates enthält und das Gewicht der Umhüllung 8 bis 12% des Kerngewichtes beträgt.

4. Formulierung nach einem der Ansprüche 1 bis 3, wobei der Kern 15 bis 25 Gewichtsteile der mikrokristallinen Zellulose enthält.

5. Formulierung nach einem der Ansprüche 1 bis 4, wobei die Umhüllung etwa 30 Gewichtsteile des hydroxylierten Zellulosederivates umfaßt.

6. Formulierung nach einem der Ansprüche 1 bis 5, wobei das hydroxylierte Zellulosederivat Hydroxypropyl-methylzellulose ist, mit einem Substitutionsgrad von 28 bis 30% der Methoxygruppen und 7 bis 12% Hydroxygruppen.

7. Formulierung nach einem der Ansprüche 1 bis 6, wobei das Akrylpolymer neutral ist.

8. Formulierung nach einem der Ansprüche 1 bis 7, wobei die Körnchen in einer Kapsel enthalten sind.

9. Verfahren zur Herstellung einer Formulierung nach einem der Ansprüche 1 bis 8, welches:

1. das Mischen von Ibuprofen mit der mikrokri-

stallinen Zellulose;

2. das Formen des Gemisches zu Teilchen;

3. das Bilden einer Lösung des hydroxylierten Zellulosederivates und des Akrylpolymeren; und

4. die Umhüllung der Teilchen unter Verwendung der Lösung zur Bildung der Körnchen umfaßt.

10. Verfahren nach Anspruch 9, wobei die Körnchen nach der Bildung gesiebt werden.

Revendications

1. Formulation d'acide α -methyl-4-(2-méthylpropyl)benzène-acétique (Ibuprofen) à libération prolongée comprenant suffisamment de granules pour former une dose d'Ibuprofen prédéterminée ou un nombre de doses d'Ibuprofen prédéterminé, chaque granule ayant un diamètre situé entre 0,5 et 2,0 mm et comprenant:

a) un noyau contenant 100 parties en poids d'Ibuprofen et de 10 à 70 parties en poids d'une cellulose microcristalline; et

b) un revêtement recouvrant sensiblement toute la surface du noyau et comprenant 100 parties en poids d'un polymère acrylique insoluble dans l'eau mais gonflable dans l'eau et 20 à 70 parties en poids d'un dérivé cellulosique hydroxylé hydrosoluble, le poids du revêtement représentant 5 à 20% du poids du noyau.

2. Formulation selon la revendication 1, qui est une formulation de 24 heures, dans laquelle le diamètre des granules est compris entre 1,5 et 2,0 mm, le revêtement contient 25 à 35 parties en poids de dérivé cellulosique hydroxylé hydrosoluble et le poids du revêtement est d'approximativement 20% du poids du noyau.

3. Formulation selon la revendication 1, qui est une formulation de 12 heures, dans laquelle le diamètre des granules est compris entre 0,7 et 1,2 mm, le revêtement contient 25 à 35 parties en poids de dérivé cellulosique hydroxylé hydrosoluble et le poids du revêtement est compris entre 8 et 12% du poids du noyau.

4. Formulation selon l'une quelconque des revendications 1 à 3, dans laquelle le noyau contient entre 15 et 25 parties en poids de cellulose microcristalline.

5. Formulation selon l'une quelconque des revendications 1 à 4, dans laquelle le revêtement comprend environ 30 parties en poids de dérivé cellulosique hydroxylé.

6. Formulation selon l'une quelconque des revendications 1 à 5, dans laquelle le dérivé cellulosique hydroxylé est une hydroxypropylmethylcellulose ayant un degré de substitution de 28 à 30% de groupes méthoxy et de 7 à 12% de groupes hydroxy.

7. Formulation selon l'une quelconque des revendications 1 à 6, dans laquelle le polymère acrylique est neutre.

8. Formulation selon l'une quelconque des revendications 1 à 7, dans laquelle les granules sont contenus dans une capsule.

9. Procédé de préparation d'une formulation selon l'une quelconque des revendications 1 à 8,

comprenant:

1. le mélange de l'ibuprofen avec la cellulose microcristalline;
2. la transformation d mélange en particules;
3. la formation d'une solution du dérivé cellulosique hydroxylé et du polymère acrylique; et

4. le revêtement des particules avec la solution pour former les granules.

10. Procédé selon la revendication 9, dans lequel les granules sont tamisés après leur formation.

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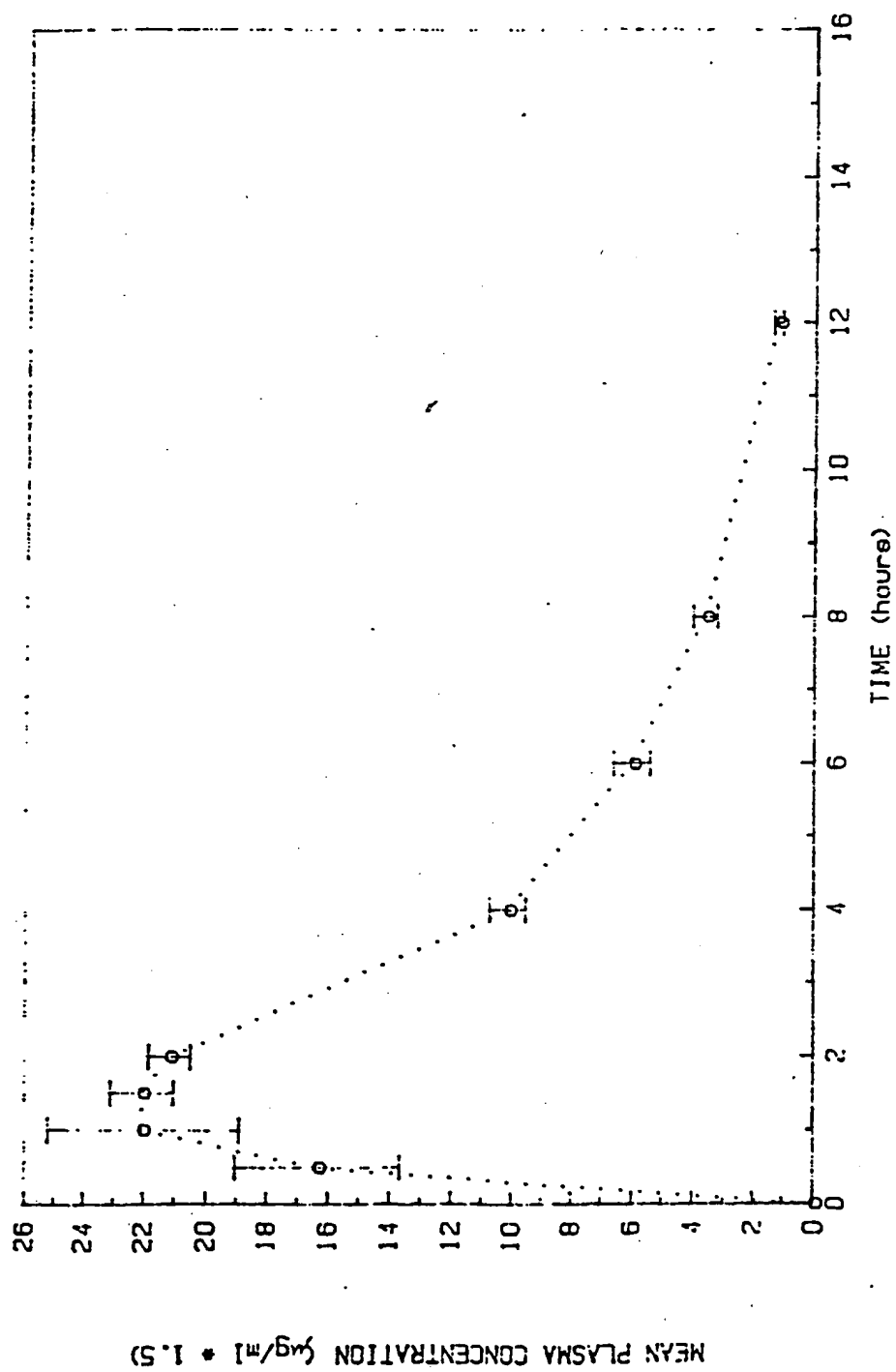
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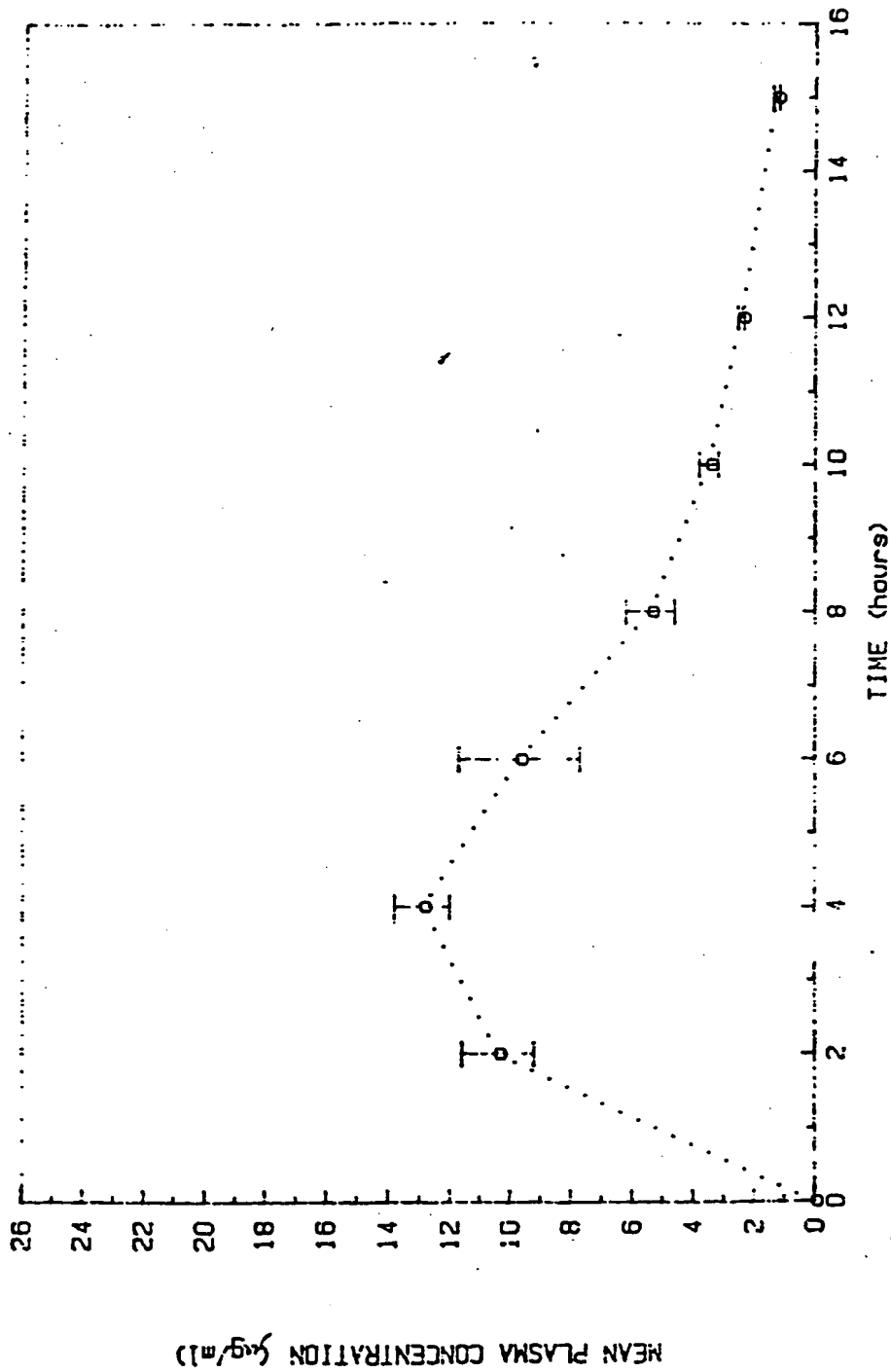
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Figure 1

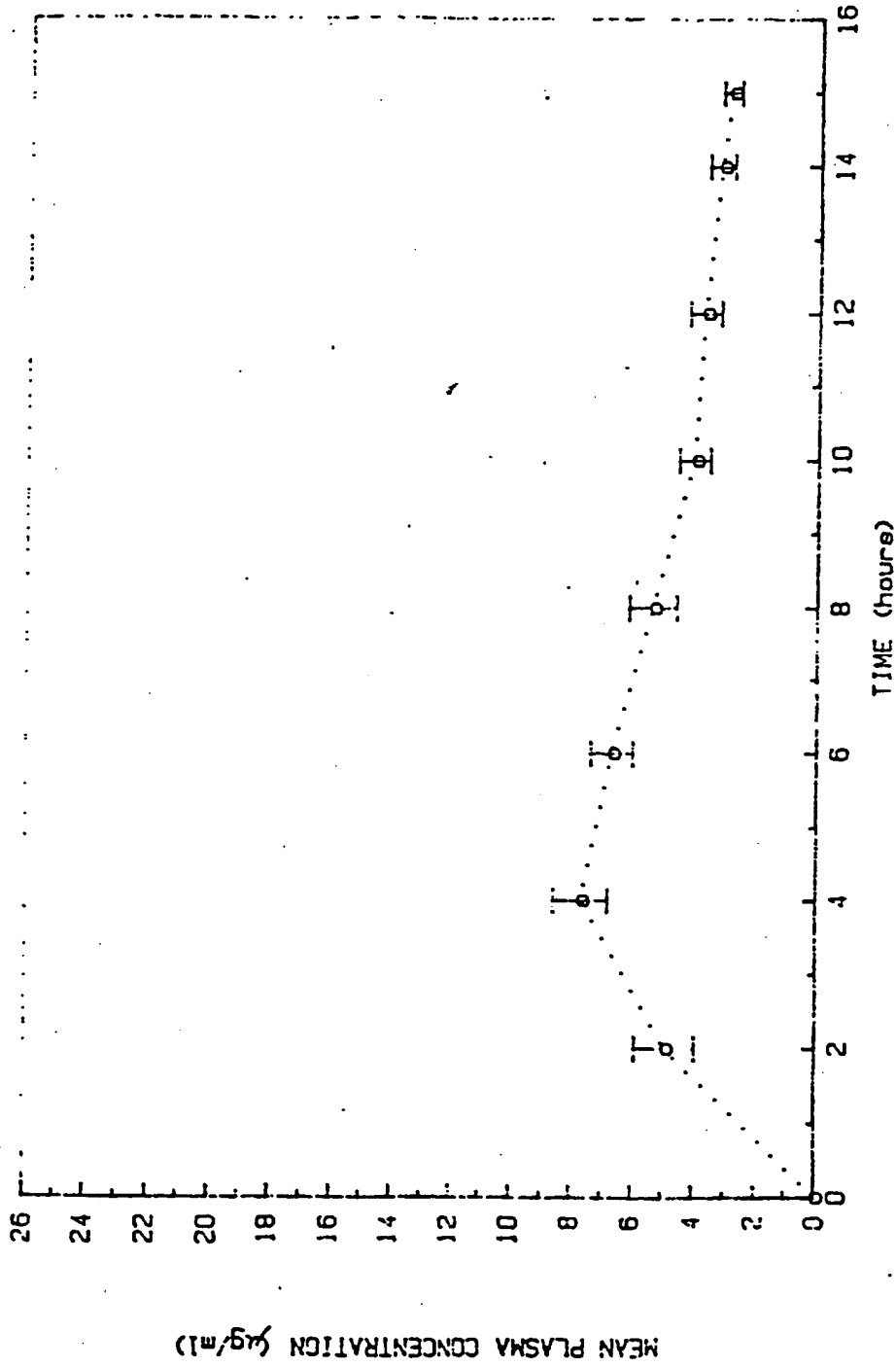


Plasma levels following administration of Brufen
(Ibuprofen 200 mg). Mean of five subjects.

Figure 2

Plasma levels (\pm SEM) following administration of
Fenbid (Ibuprofen 300 mg). Mean of five subjects.

Figure 3



Plasma levels (±SEM) following administration of
Product II 300 mg. Mean of five subjects.

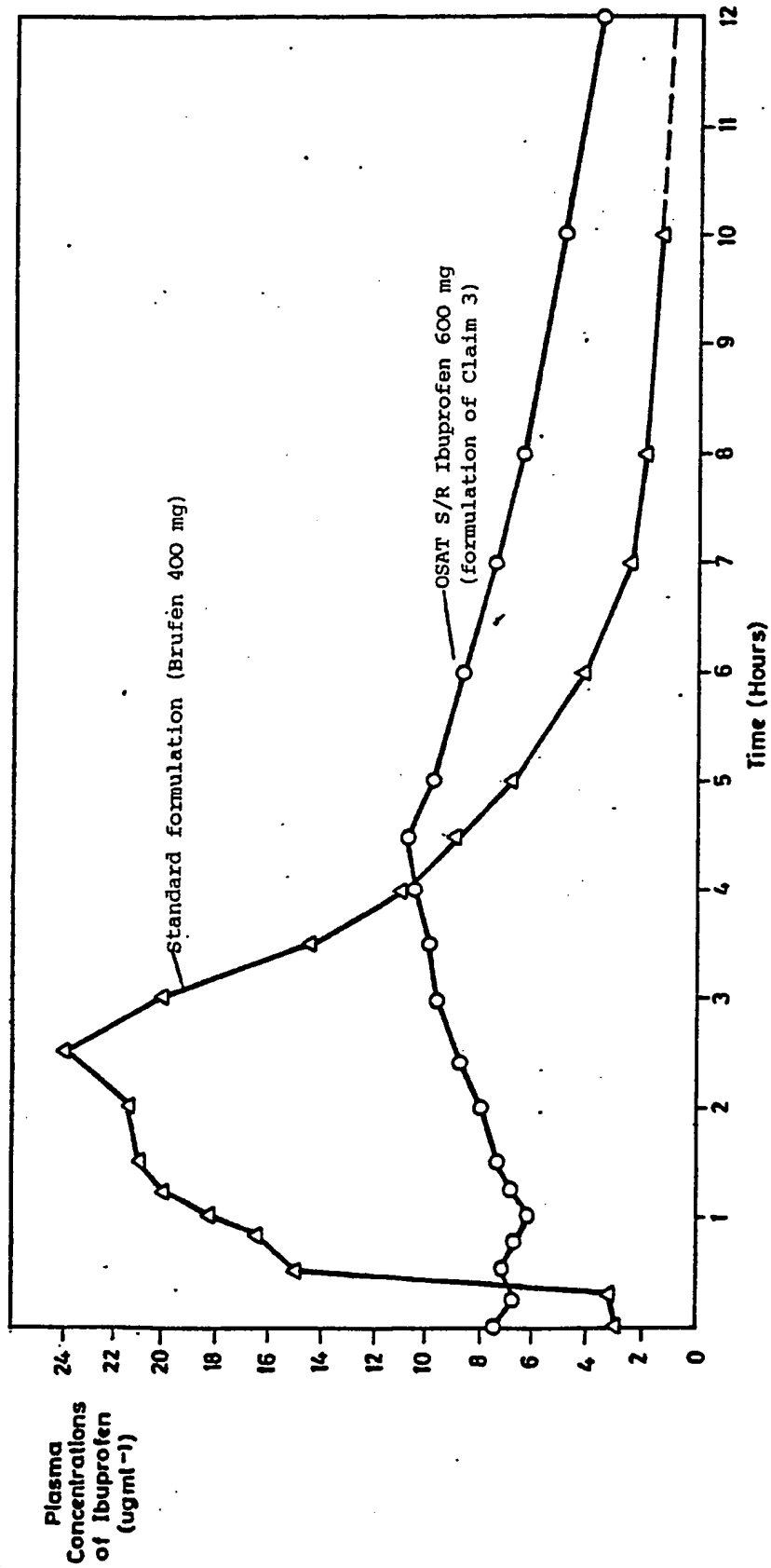


FIG 4. Blood level time profiles after a fourth day morning dose following three days of OSAT S/R Ibuprofen (600mg b.d.) and a standard formulation (Brufen 400 mg t.i.d.)